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		FIRST NAMED INVENTOR	ATTORNEY DOCKET NO.	CONFIRMATION NO.	
APPLICATION NO.	FILING DATE	FIRST WANTED IN VENTOR			
09/785,514	02/16/2001	Jian-Bing Fan	A-689/0-1/DJB/KM5/DCF	3302	
7:	590 04/15/2002				
	NNO. FILING DATE FIRST NAMED INVENTOR ATTORIST STATES A STATEST ALBRITTON & HERBERT LLP CHAKRABARTI, ARUN K mbarcadero Center, Suite 3400				
FLEHR HOHBACH TEST ALBRITTON & HERBERT LLP Four Embarcadero Center, Suite 3400 San Francisco, CA 94111-4187			CHAKRABARTI, ARUN K		
			ART UNIT	PAPER NUMBER	
			1634		
	DATE MAILED: 04/15/2002				

Please find below and/or attached an Office communication concerning this application or proceeding.

<u> </u>		Application No.		Applicant(s)				
Office Action Summary		09/785,514		FAN ET AL.				
		Examiner		Art Unit				
		Diana Johannse		1634				
	- The MAILING DATE of this communication app	pears on the cove	r sheet with the c	orrespondence a	ddress			
Period fo		VIO SET TO EV	DIDE 2 MONTH/	S) FROM				
THE N - Extending after S - If the - If NO - Failuring Any re	ORTENED STATUTORY PERIOD FOR REPLY MAILING DATE OF THIS COMMUNICATION. Issions of time may be available under the provisions of 37 CFR 1.1 SIX (6) MONTHS from the mailing date of this communication. It period for reply specified above is less than thirty (30) days, a reply period for reply is specified above, the maximum statutory period are to reply within the set or extended period for reply will, by statute eply received by the Office later than three months after the mailing and patent term adjustment. See 37 CFR 1.704(b).	136(a). In no event, how ly within the statutory mi will apply and will expire	ever, may a reply be tin nimum of thirty (30) day SIX (6) MONTHS from to become ABANDONE	nely filed s will be considered time the mailing date of this (35 U.S.C. § 133).	ely. communication.			
1)⊠	Responsive to communication(s) filed on 25	March 2002 .						
2a) ☐	This action is FINAL . 2b)⊠ Th	his action is non-						
3)	Since this application is in condition for allow closed in accordance with the practice under	vance except for to r Ex parte Quayle	formal matters, p e, 1935 C.D. 11,	rosecution as to 453 O.G. 213.	the merits is			
•	ion of Claims	on .						
	 Claim(s) 1-26 is/are pending in the application. 4a) Of the above claim(s) 1-13 is/are withdrawn from consideration. 							
	Claim(s) is/are allowed.							
·	Claim(s) <u>14-26</u> is/are rejected.							
	Claim(s) is/are objected to. Claim(s) are subject to restriction and/	or election requir	ement.					
	tion Papers	-						
• -	The specification is objected to by the Examin	ner.						
10)□	The drawing(s) filed on is/are: a) ☐ acc	epted or b) dobje	cted to by the Ex	aminer.				
	Applicant may not request that any objection to t	the drawing(s) be h	eld in abeyance.	See 37 CFR 1.85(a	à).			
11)	The proposed drawing correction filed on	is: a)☐ appro	ved b)⊡ disapp	roved by the Exan	niner.			
	If approved, corrected drawings are required in r		action.					
12)	The oath or declaration is objected to by the E	Examiner.						
Priority	under 35 U.S.C. §§ 119 and 120							
13)	Acknowledgment is made of a claim for forei	ign priority under	35 U.S.C. § 119	(a)-(d) or (f).				
а) All b) Some * c) None of:							
	1. Certified copies of the priority documents have been received.							
	2. Certified copies of the priority documents have been received in Application No							
	3. Copies of the certified copies of the prapplication from the International I See the attached detailed Office action for a li	ist of the certified	copies not recei	ived.				
14)	Acknowledgment is made of a claim for dome	estic priority unde	r 35 U.S.C. § 11	9(e) (to a provision	nal application).			
	a) The translation of the foreign language Acknowledgment is made of a claim for dome	provisional applic	ation has been r	received.				
Attachm								
2) \ \ \ \ \ \ \ \ \ \ \ \ \ \ \ \ \ \ \	otice of References Cited (PTO-892) otice of Draftsperson's Patent Drawing Review (PTO-948) formation Disclosure Statement(s) (PTO-1449) Paper No(s	5) s)	☐ Interview Sumn☐ Notice of Inform☐ Other: Detailed	nary (PTO-413) Pape nal Patent Application I Action .	r No(s) (PTO-152)			
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Art Unit: 1634

DETAILED ACTION

Specification

1. Applicant's election of claims 14-26, corresponding to Group II, in paper Number 11 is hereby acknowledged. Non-elected claims 1-13 have been canceled without prejudice towards further prosecution.

Claim 21 is misnumbered. It has two steps having the same number c). Proper correction is advised.

Double Patenting

2. The nonstatutory double patenting rejection is based on a judicially created doctrine grounded in public policy (a policy reflected in the statute) so as to prevent the unjustified or improper timewise extension of the "right to exclude" granted by a patent and to prevent possible harassment by multiple assignees. See *In re Goodman*, 11 F.3d 1046, 29 USPQ2d 2010 (Fed. Cir. 1993); *In re Longi*, 759 F.2d 887, 225 USPQ 645 (Fed. Cir. 1985); *In re Van Ornum*, 686 F.2d 937, 214 USPQ 761 (CCPA 1982); *In re Vogel*, 422 F.2d 438, 164 USPQ 619 (CCPA 1970);and, *In re Thorington*, 418 F.2d 528, 163 USPQ 644 (CCPA 1969).

A timely filed terminal disclaimer in compliance with 37 CAR 1.321© may be used to overcome an actual or provisional rejection based on a nonstatutory double patenting ground provided the conflicting application or patent is shown to be commonly owned with this application. See 37 CAR 1.130(b).

Effective January 1, 1994, a registered attorney or agent of record may sign a terminal disclaimer. A terminal disclaimer signed by the assignee must fully comply with 37 CAR 3.73(b).

3. Claims 14-26 are rejected under the judicially created doctrine of obviousness-type double patenting as being unpatentable over claims 1-30 of U.S. Patent No. 6,355,431 B1.

Page 3

Art Unit:

Although the conflicting claims are not identical, they are not patentably distinct from each other because claims 1-30 of U.S. Patent No. 6,355,431 B1 clearly teach the instant claimed method of genotyping comprising:

- a) providing an array composition comprising:
 - I) a substrate with a surface comprising discrete sites; and
- ii) a population of microspheres comprising at least a first and a second subpopulation, wherein the microspheres of each subpopulation each comprise a plurality of target analytes attached to the microspheres with first and second attachment moieties, respectively;

wherein the microspheres are randomly distributed on the surface;

- b) contacting the array composition with a first set of extension probes that hybridize with at least the first target sequence adjacent to a first detection position to form an extension complex;
 - c) contacting the extension complex with a composition comprising:
 - I) at least a first nucleotide;
 - ii) polymerase;

wherein the polymerase extends a first extension probe with the first nucleotide when the first nucleotide is complementary to the first detection position of the first target sequence; and

d) detecting the presence of a first nucleotide and also a method further comprising adding a ligase to form a ligation complex (Claim 6).

Art Unit:

Claim Rejections - 35 USC § 102

4. The following is a quotation of the appropriate paragraphs of 35 U.S.C. 102 that form the basis for the rejections under this section made in this Office action:

A person shall be entitled to a patent unless --

- (e) the invention was described in a patent granted on an application for patent by another filed in the United States before the invention thereof by the applicant for patent, or on an international application by another who has fulfilled the requirements of paragraphs (1), (2), and (4) of section 371© of this title before the invention thereof by the applicant for patent.
- 5. Claims 14-26 are rejected under 35 U.S.C. 102 (e) as being anticipated by Beattie et al. (U.S. Patent 6,268,147 B1) (July 31, 2001).

Beattie et al teach a method comprising:

- a) providing an array composition (Figure 6, Column 6, lines 6-13, and Example 18) comprising:
- I) a substrate with a surface comprising discrete sites (Figures 6, 15A-B); and
- ii) a population of microspheres comprising at least a first and a second subpopulation, wherein the microspheres of each subpopulation each comprise a plurality of target analytes (Figures 15A-B and Example 18);

wherein the microspheres are randomly distributed on the surface (Example 18);

b) contacting the array composition with a first set of readout probes (Figures 13 A-B and Example 18);

Art Unit:

- c) detecting the presence of a first target analyte (Figures 13 A-B and Example 18);
- d) contacting the array composition with a second set of readout probes (Figures 13 A-B and Example 18);
- e) detecting the presence of a second target analyte (Example 18), wherein the two sets of probes comprise two sets of labels (Example 18).

Beattie et al teach a method further comprising detecting the first label as an indication of the presence of the first target analyte (Example 18).

Beattie et al teach a method, wherein the first and second target sources are first and second patients (Figure 4, and Example 2).

Beattie et al inherently teach a method of genotyping comprising:

- a) providing an array composition (Figure 6, Column 6, lines 6-13, and Example 18) comprising:
- I) a substrate with a surface comprising discrete sites (Figures 6, 15A-B); and
- ii) a population of microspheres comprising at least a first and a second subpopulation, wherein the microspheres of each subpopulation each comprise a plurality of target analytes attached to the microspheres with first and second attachment moieties, respectively (Figures 15A-B and Example 18);

wherein the microspheres are randomly distributed on the surface ();

Art Unit:

and

b) contacting the array composition with a first set of extension probes that hybridize with at least the first target sequence adjacent to a first detection position to form an extension complex (Figure 14);

- c) contacting the extension complex with a composition comprising:
- I) at least a first nucleotide (Figure 14);
- ii) polymerase (Example 12 and Figure 14);

wherein the polymerase extends a first extension probe with the first nucleotide when the first nucleotide is complementary to the first detection position of the first target sequence (Example 12); and

d) detecting the presence of a first nucleotide (Example 18).

Beattie et al inherently teach a method, wherein the first nucleotide comprises a label (Example 18).

Beattie et al inherently teach a method of determining the identification of a nucleotide at a detection position in at least a first target sequence comprising:

- a) providing an array composition (Figure 6, Column 6, lines 6-13, and Example 18)) comprising:
 - I) a substrate with a surface comprising discrete sites (Figures 6, 15A-B);

Art Unit:

ii) a population of microspheres comprising at least a first and a second subpopulation, wherein the microspheres of each subpopulation each comprise a plurality of target analytes (Figures 15A-B and Example 18);

- b) forming a first hybridization complex between the target sequence and at least a first readout probe (Example 18);
 - c) determining the nucleotide at the detection position (Example 18).

Beattie et al teach a method further comprising adding a ligase to form a ligation complex (Column 12, line 48 to column 13, line 13).

Conclusion

6. Any inquiry concerning this communication or earlier communications from the examiner should be directed to Arun Chakrabarti, Ph.D. whose telephone number is (703) 306-5818.

If attempts to reach the examiner by telephone are unsuccessful, the examiner's supervisor, W. Gary Jones, can be reached on (703) 308-1152. Any inquiry of a general nature or relating to the status of this application should be directed to the Group analyst Chantae Dessau whose telephone number is (703) 605-1237. Papers related to this application may be submitted to Technology Center 1600 by facsimile transmission via the P.T.O. Fax Center located In Crystal Mall 1. The CM1 Fax Center numbers for Technology Center 1600 are either (703) 305-3014 or (703) 308-4242. Please note that the faxing of such papers must conform with the Notice to Comply published In the Official Gazette, 1096 OG 30 (November 15, 1989).

Art Unit:

Arun Chakrabarti Patent Examiner Art Unit 1634

April 8, 2002

ARUN K. CHAKRABART PATENT EXAMINER